WHAT IS CLAIMED IS:

A compound of formula I

$$R_2$$
 $(CR_3R_4)n$
 $Q-R_5$
 (I)

5 wherein

20

Q is SO₂, CO, CONR₂₄, CSNR₂₅ or CH₂;

W is N or CR6;

X is N or CR,;

Y is NR₈ or CR₉R₁₀;

n is 0 or an integer of 1 or 2;

Z is NR_{11} or $CR_{12}R_{13}$ with the proviso that when n is 1, Q is SO_2 CO or CH_2 and W is CR_6 then Z must be $CR_{12}R_{13}$ and with the further provisos that when Y is NR_8 then Z must be $CR_{12}R_{13}$ and at least one of Y and Z

15 must be NR_8 or NR_{11} ;

 $\rm R_1$, $\rm R_2$ and $\rm R_7$ are each independently H, halogen, CN, ${\rm OCO_2R_{14},\ CO_2R_{15},\ CONR_{29}R_{30},\ CNR_{16}NR_{17}R_{18},\ SO_mR_{19},\ NR_{20}R_{21},}\\ {\rm OR_{22},\ COR_{23}\ or\ a\ C_1-C_6alkyl,\ C_2-C_6alkenyl,\ C_2-C_6alkynyl,}\\ {\rm C_3-C_6cycloalkyl,\ cycloheteroalkyl,\ aryl\ or\ heteroaryl\ group\ each\ optionally\ substituted;}$

 R_3 , R_4 , R_9 , R_{10} , R_{12} and R_{13} are each independently H or an optionally substituted C_1-C_6 alkyl group;

 R_s is an optionally substituted C_1-C_6 alkyl, aryl or heteroaryl group;

m is 0 or an integer of 1 or 2;

 R_6 is H, halogen, or an optionally substituted C_1 - C_6 alkyl, C_1 - C_6 alkoxy, aryl or heteroaryl group;

- R_s and R_{11} are each independently H, $CNR_{26}NR_{27}R_{28}$ or a C_1 C_6 alkyl, C_3 - C_6 cycloalkyl, cycloheteralkyl, aryl or heteroaryl group each optionally substituted;
- R_{14} , R_{15} , R_{22} and R_{23} are each independently H or an optionally substituted C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_3 - C_6 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group;
- R_{16} , R_{17} , R_{18} , R_{20} , R_{21} , R_{26} , R_{27} , R_{28} , R_{29} and R_{30} are each independently H or C_1-C_4 alkyl;
- 10 R_{19} is an optionally substituted C_1-C_6 alkyl, aryl or heteroaryl group;
 - $\rm R_{24}$ and $\rm R_{25}$ are each independently H or an optionally substituted $\rm C_1-C_6 alkyl,$ aryl or heteroaryl group; and
- 15 ____ represents a single bond or a double bond; or the stereoisomers thereof or the pharmaceutically acceptable salts thereof.
- $\hbox{2.} \quad \hbox{The compound according to claim 1 wherein Y is} \\ \hbox{20} \quad \hbox{NR}_{_{\!R}}.$
 - 3. The compound according to claim 1 wherein n is 0 or 2.
- 25 4. The compound according to claim 1 wherein W is N.
 - 5. The compound according to claim 2 wherein n is 1.
- $\mbox{6.} \quad \mbox{The compound according to claim 4 wherein Z is } \\ 30 \quad \mbox{NR}_{11}.$
 - 7. The compound according to claim 5 wherein Q is SO_2 and $R_{\mbox{\tiny 5}}$ is an optionally substituted aryl or heteroaryl group.

- 8. The compound according to claim 7 wherein X is CH and ____ represents a single bond.
- 9. The compound according to claim 1 selected from 5 the group consisting of:
 - 1-(phenylsulfonyl)-3-(piperidin-4-yl)-1H-indazole;
 - 1-(4-nitrophenyl)-3-(piperidin-4-yl)-1H-indazole;
 - 1-(4-fluorophenyl)-3-(piperidin-4-yl)-1H-indazole;
 - 1-(3,4-dimethoxyphenyl-3-(piperidin-4-yl)-1H-indazole;
- 10 1-(4-fluorophenylsulfonyl)-3-(1-methyl-pyrrolidin-3-yl)-1Hindole;
 - 1-(4-chlorophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-indole;
 - 1-(naphth-2-ylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-indole;
 - 1-(4-aminophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-indole;
 - 1-(3,4-dimethoxyphenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-indole;
- 20 1-(3,4-dichlorophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-indole;
 - 1-[(4,5-dichlorothien-2-yl)sulfonyl]-3-(1-methyl-pyrrolidin-3-yl)-1H-indole;
 - 1-(2-bromophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-indole;
 - 1-(4-iodophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1Hindole;
 - 1-(2-iodophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1Hindole;
- 30 1-(4-aminophenylsulfonyl)-3-(1-benzylpyrrolidin-3-yl)-1Hindole;
 - 3-(1-benzylpyrrolidin-3-yl)-1-(4-methylphenylsulfonyl)-1H-indole;
- 3-(1-benzylpyrrolidin-3-yl)-1-(3,4-dichlorophenyl-sulfonyl)35 1H-indole;

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3-(1-benzylpyrrolidin-3-yl)-1-(2-bromophenylsulfonyl)-1H-
        indole;
   5-[3-(1-benzylpyrrolidin-3-yl)-indole-1-sulfonyl]-4-methyl-
        thiazol-2-ylamine;
   3-(1-benzylpyrrolidin-3-yl)-1-[(5-bromothien-2-yl)sulfonyl]-
        1H-indole;
   1-phenylsulfonyl-3-(1-methylpyrrolidin-3-yl)-1H-pyrrolo[2,3-
        b]pyridine;
   1-phenylsulfonyl-3-(1-methylpyrrolidin-3-yl)-1H-indazole;
   1-phenylsulfonyl-3-(1-methyl-2,5-dihydro-1H-pyrrol-3-yl)-1H-
        pyrrolo[2,3-b]pyridine;
   1-phenylsulfonyl-3-(1-methyl-2,5-dihydro-1H-pyrrol-3-yl)-1H-
        indole;
    1-phenylsulfonyl-3-(1-methylpiperidin-4-yl)-1H-indazole;
   1-phenylsulfonyl-3-(1-methyl-1,2,3,6-tetrahydropyridin-4-
        yl)-1H-indazole;
    1-phenylsulfonyl-3-(1-methylazepan-4-yl)-1H-pyrrolo[2,3-
        b]pyridine;
    1-phenylsulfonyl-3-(1-methylazepan-4-yl)-1H-indole;
    1-phenylsulfonyl-5-fluoro-3-(1-methylazepan-4-yl)-1H-indole;
    1-phenylsulfonyl-3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-
         yl)-1H-indole;
    1-phenylsulfonyl-3-(1-methyl-2,5,6,7-tetrahydro-1H-azepin-4-
         yl)-1H-indole;
    1-phenylsulfonyl-3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-
25
         yl)-1H-pyrrolo[2,3-b]pyridine;
    1-phenylsulfonyl-5-fluoro-3-(1-methyl-2,3,6,7-tetrahydro-1H-
         azepin-4-yl)-1H-indole;
    1-phenylsulfonyl-5-fluoro-3-(1-methyl-2,5,6,7-tetrahydro-1H-
         azepin-4-y1)-1H-indole;
30
    1-(benzo[b]thioen-4-ylsulfonyl)-3-(1-methyl-pyrrolidin-3-
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1-(3-fluorophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-

y1)-1H-pyrrolo[2,3-b]pyridine;

indazole;

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1-(2,5-dichlorophenylsulfonyl)-3-(2,5-dihydro-1H-pyrrol-3-
        yl)-1H-pyrrolo[2,3-b]pyridine;
   8-[3-(1-methyl-2,5-dihydro-1H-pyrrol-3-yl)indole-1-
         sulfonyl]-quinoline;
   1-phenylsulfonyl-5-chloro-3-(1-methylpiperidin-4-yl)-1H-
5
         indazole;
    5-methoxy-3-(1-methyl-1,2,3,6-tetrahydropyridin-4-yl)-1-
         (naphth-1-yl-sulfonyl)-1H-indazole;
    3-(1-methylazepan-4-yl)-1-(naphth-1-yl-sulfonyl)-1H-
         pyrrolo[2,3-b]pyridine;
10
    3-(1-methylazepan-4-yl)-1-(naphth-1-yl-sulfonyl)-1H-indole;
    1-(benzo[b]thien-4-ylsulfonyl)-5-fluoro-3-(1-methylazepan-4-
         yl)-1H-indole;
    8-[3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-yl)-indole-1-
         sulfonyl]-quinoline;
15
    3-(1-methyl-2,5,6,7-tetrahydro-1H-azepin-4-yl)-1-(naphth-1-
         ylsulfonyl)-1H-indole;
    8-[3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-yl)-
         pyrrolo[2,3-b]pyridine-1-sulfonyl]-quinoline;
    8-[5-fluoro-3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-yl)-
20
         indole-1-sulfonyl]-quinoline;
    5-fluoro-3-(1-methyl-2,5,6,7-tetrahydro-1H-azepin-4-yl)-1-
         (naphth-1-ylsulfonyl)-1H-indole;
    1-(benzo[b]thien-4-ylsulfonyl)-3-(1-benzyl-pyrrolidin-3-yl)-
         1H-pyrrolo[2,3-b]pyridine;
25
    1-(3-fluoro-phenylsulfonyl)-3-(1-phenethyl-pyrrolidin-3-yl)-
         1H-indazole;
    1-(2,5-dichlorophenylsulfonyl)-3-(1-ethyl-2,5-dihydro-1H-
         pyrrol-3-yl)-1H-pyrrolo[2,3-b]pyridine;
    3-(1-methyl-2,5-dihydro-1H-pyrrol-3-yl)-1-(naphth-2-
30
         ylsulfonyl)-1H-indole;
    5-chloro-1-(3-fluorophenylsulfonyl)-3-piperidin-4-yl-1H-
          indazole;
     5-methoxy-1-(naphth-1-ylsulfonyl)-3-(1,2,2-trimethyl-
          1,2,3,6-tetrahydro-pyridin-4-yl)-1H-indazole;
35
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1-(naphth-1-ylsulfonyl)-3-(1-phenethyl-azepan-4-yl)-1H-
        pyrrolo[2,3-b]pyridine;
   3-azepan-4-yl-1-(naphth-1-ylsulfonyl)-1H-indole;
   3-azepan-4-yl-1-(3-chloro-5-methyl-benzo[b]thien-2-
         vlsulfonyl)-5-fluoro-1H-indole;
5
    8-[3-(1-phenethyl-2,3,6,7-tetrahydro-1H-azepin-4-yl)-indole-
         1-sulfonyl]-quinoline;
   3-[1-(3,3-dimethylbutyl)-2,5,6,7-tetrahydro-1H-azepin-4-yl]-
         1-(naphth-2-ylsulfonyl)-1H-indole;
    1-(2,3-dichlorophenylsulfonyl)-3-(1-methyl-2,3,6,7-
10
         tetrahydro-1H-azepin-4-yl)-1H-pyrrolo[2,3-]pyridine;
    1-[(3-chloro-5-methoxyphenylsulfonyl)]-3-(2,2-dimethyl-
         2,3,6,7-tetrahydro-1H-azepin-4-yl)-5-fluoro-1H-indole;
    3-azepan-4-yl-5-fluoro-1-(naphth-2-ylsulfonyl)-1H-indole;
    1-Benzenesulfonyl-3-piperidin-3-yl-1H-indole;
15
    1-(4-isopropyl-benzenesulfonyl)-3-piperidin-3-yl-1H-indole;
    1-(5-chloro-thiophene-2-sulfonyl)-3-piperidin-3-yl-1H-
         indole;
    1-(3-chloro-benzenesulfonyl)-3-piperidin-3-yl-1H-indole;
    1-(3,4-difluoro-benzenesulfonyl)-3-piperidin-3-yl-1H-indole;
20
    1-(4-trifluoromethoxy-benzenesulfonyl)-3-piperidin-3-yl-1H-
         indole;
    1-(4-methoxy-benzenesulfonyl)-3-piperidin-3-yl-1H-indole;
    1-(4-trifluoromethy-benzenesulfonyl)-3-piperidin-3-yl-1H-
25
         indole;
    1-(3-chloro-4-methyl-benzenesulfonyl)-3-piperidin-3-yl-1H-
         indole;
    1-(2-chloro-4-trifluoromethyl-benzenesulfonyl)-3-piperidin-
         3-yl-1H-indole;
    1-(2-naphthylenesulfonyl)-3-piperidin-3-yl-1H-indole;
30
    1-(5-chloro-3-methyl-benzo[b]thiophene-2-sulfonyl)-3-
         piperidin-3-yl-1H-indole;
    1-(2,6-dichloro-imidazo[2,1-b]thiazole-5-sulfonyl)-3-
         piperidin-3-yl-1H-indole;
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- 2-chloro-3-(3-piperidin-3-yl-indole-1-sulfonyl)-imidazo[1,2-a]pyridine;
- 2-chloro-3-(3-piperidin-3-yl-indole-1-sulfonyl)benzo[d]imidazo[2,1-b]thiazole;
- 5 1-(4-isopropyl-benzenesulfonyl)-3-piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
 - 1-(5-chloro-thiophene-2-sulfonyl)-3-piperidin-3-yl-1Hpyrrolo[2,3-b]pyridine;
 - 1-(3-chloro-benzenesulfonyl)-3-piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
 - 1-(3,4-difluoro -benzenesulfonyl)-3-piperidin-3-yl-1Hpyrrolo[2,3-b]pyridine;
 - 1-(4-trifluoromethoxy-benzenesulfonyl)-3-piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
- 15 1-(3-chloro-4-methyl-benzenesulfonyl)-3-piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
 - 1-(2-chloro-4-trifluoromethyl-benzenesulfonyl)-3-piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
 - 1-(2-naphthylenesulfonyl)-3- piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
 - 1-(5-chloro-3-methyl-benzo[b]thiophene-2-sulfonyl)-3-piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
 - 2-chloro-3-(3-piperidin-3-yl-pyrrolo[2,3-b]pyridine-1-sulfonyl)-imidazo[1,2-a]pyridine;
- 25 2-chloro-3-(3-piperidin-3-yl-pyrrolo[2,3-b]pyridine-1-sulfonyl)-benzo[d]imidazo[2,1-b]thiazole; and the pharmaceutically acceptable salts thereof.
- 10. A method for the treatment of a disorder of the
 30 central nervous system related to or affected by the 5-HT6
 receptor in a patient in need thereof which comprises
 providing said patient with a therapeutically effective
 amount of a compound of formula I

$$R_2$$
 $(CR_3R_4)n$
 (CR_3R_5)
 (I)

wherein

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Q is SO_2 , CO, $CONR_{24}$, $CSNR_{25}$ or CH_2 ;

W is N or CR₆;

X is N or CR,;

Y is NR, or CR,R10;

n is 0 or an integer of 1 or 2;

Z is NR_{11} or $CR_{12}R_{13}$ with the proviso that when n is 1, Q is SO_2 , CO or CH_2 and W is CR_6 then Z must be $CR_{12}R_{13}$ and with the further provisos that when Y is NR_8 then Z must be $CR_{12}R_{13}$ and at least one of Y and Z must be NR_8 or NR_{11} ;

 R_1 , R_2 and R_7 are each independently H, halogen, CN, OCO_2R_{14} , CO_2R_{15} , $CONR_{29}R_{30}$, $CNR_{16}NR_{17}R_{18}$, SO_mR_{19} , $NR_{20}R_{21}$, OR_{22} , COR_{23} or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;

 R_3 , R_4 , R_9 , R_{10} , R_{12} and R_{13} are each independently H or an optionally substituted C_1-C_6 alkyl group;

 R_s is an optionally substituted C_1 - C_6 alkyl, aryl or heteroaryl group;

m is 0 or an integer of 1 or 2;

R₆ is H, halogen, or an optionally substituted C₁-C₆alkyl, C₁-C₆alkoxy, aryl or heteroaryl group;

- R_{s} and R_{11} are each independently H, $CNR_{26}NR_{27}R_{28}$ or a C_{1} C_{6} alkyl, C_{3} - C_{6} cycloalkyl, cycloheteralkyl, aryl or heteroaryl group each optionally substituted;
- R_{14} , R_{15} , R_{22} and R_{23} are each independently H or an optionally substituted C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_3 - C_6 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group;
- R_{16} , R_{17} , R_{18} , R_{20} , R_{21} , R_{26} , R_{27} , R_{28} , R_{29} and R_{30} are each independently H or C_1 - C_4 alkyl;
- 10 R₁₉ is an optionally substituted C₁-C₆alkyl, aryl or heteroaryl group;
 - $\rm R_{24}$ and $\rm R_{25}$ are each independently H or an optionally substituted $\rm C_1-C_6 alkyl,$ aryl or heteroaryl group; and
- 15 ____ represents a single bond or a double bond; or the stereoisomers thereof or the pharmaceutically acceptable salts thereof.
- 11. The method according to claim 10 wherein said 20 disorder is a mood disorder, a motor disorder, or a cognitive disorder.
 - 12. The method according to claim 10 wherein said disorder is schizophrenia.
 - 13. The method according to claim 11 wherein said disorder is anxiety or depression.
- 14. The method according to claim 11 wherein said disorder is memory loss or attention deficit disorder.
 - 15. A pharmaceutical composition which comprises a pharmaceutically acceptable carrier and an effective amount of a compound of formula I

$$R_2$$
 $(CR_3R_4)n$
 $Q-R_5$
 (I)

wherein

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25

Q is SO2, CO, CONR24, CSNR25 or CH2;

W is N or CR,;

5 X is N or CR_7 ;

Y is NR, or CR,R10;

n is 0 or an integer of 1 or 2;

Z is NR_{11} or $CR_{12}R_{13}$ with the proviso that when n is 1, Q is SO_2 , CO or CH_2 , and W is CR_6 then Z must be $CR_{12}R_{13}$ and with the further provisos that when Y is NR_8 then Z must be $CR_{12}R_{13}$ and at least one of Y and Z must be NR_8 or NR_{11} ;

 $\rm R_1$, $\rm R_2$ and $\rm R_7$ are each independently H, halogen, CN, $\rm OCO_2R_{14}, \ CO_2R_{15}, \ CONR_{29}R_{30}, \ CNR_{16}NR_{17}R_{18}, \ SO_mR_{19}, \ NR_{20}R_{21}, \\ \rm OR_{22}, \ COR_{23} \ or \ a \ C_1-C_6alkyl, \ C_2-C_6alkenyl, \ C_2-C_6alkynyl, \\ \rm C_3-C_6cycloalkyl, \ cycloheteroalkyl, \ aryl \ or \\ \rm heteroaryl \ group \ each \ optionally \ substituted;$

 R_3 , R_4 , R_9 , R_{10} , R_{12} and R_{13} are each independently H or an optionally substituted C_1-C_6 alkyl group;

 R_s is an optionally substituted C_1-C_6 alkyl, aryl or heteroaryl group;

m is 0 or an integer of 1 or 2;

 $R_{\rm G}$ is H, halogen, or an optionally substituted $C_{\rm 1}$ - $C_{\rm G}$ alkyl, $C_{\rm 1}$ - $C_{\rm G}$ alkoxy, aryl or heteroaryl group;

 R_{s} and R_{11} are each independently H, $CNR_{26}NR_{27}R_{28}$ or a C_{1} - C_{6} alkyl, C_{3} - C_{6} cycloalkyl, cycloheteralkyl, aryl or heteroaryl group each optionally substituted;

10

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- R₁₄, R₁₅, R₂₂ and R₂₃ are each independently H or an optionally substituted C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group;
- R_{16} , R_{17} , R_{18} , R_{20} , R_{21} , R_{26} , R_{27} , R_{28} , R_{29} and R_{30} are each independently H or C_1-C_4 alkyl;
 - R₁₉ is an optionally substituted C₁-C₆alkyl, aryl or heteroaryl group;
- R_{24} and R_{25} are each independently H or an optionally substituted C_1-C_6 alkyl, aryl or heteroaryl group; and
- --- represents a single bond or a double bond; or the stereoisomers thereof or the pharmaceutically acceptable salts thereof.
- 16. The composition according to claim 15 having a formula I compound wherein n is 1; Q is SO_2 ; Y is NR_8 ; and X is CR_2 .
- 20 17. The composition according to claim 15 having a formula I compound wherein n is 0; Q is SO_2 ; X is CR_7 ; and Z is NR_{11} .
- 18. The composition according to claim 16 having a formula I compound wherein $R_{\rm s}$ is an optionally substituted aryl group and --- represents a single bond.
 - 19. The composition according to claim 15 having a formula I compound selected from the group consisting of:
- 30 1-(phenylsulfonyl)-3-(piperidin-4-yl)-1H-indazole;
 - 1-(4-nitrophenyl)-3-(piperidin-4-yl)-1H-indazole;
 - 1-(4-fluorophenyl)-3-(piperidin-4-yl)-1H-indazole;
 - 1-(3,4-dimethoxyphenyl-3-(piperidin-4-yl)-1H-indazole;
 - 1-(4-fluorophenylsulfonyl)-3-(1-methyl-pyrrolidin-3-yl)-1Hindole;

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1-(4-chlorophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-
         indole;
   1-(naphth-2-ylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-
         indole;
   1-(4-aminophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-
5
         indole;
    1-(3,4-dimethoxyphenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-
         1H-indole;
   1-(3,4-dichlorophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-
10
         1H-indole;
   1-[(4,5-dichlorothien-2-yl)sulfonyl]-3-(1-methyl-pyrrolidin-
         3-y1)-1H-indole;
    1-(2-bromophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-
         indole;
    1-(4-iodophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-
15
         indole;
    1-(2-iodophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-
         indole:
    1-(4-aminophenylsulfonyl)-3-(1-benzylpyrrolidin-3-yl)-1H-
20
         indole;
    3-(1-benzylpyrrolidin-3-yl)-1-(4-methylphenylsulfonyl)-1H-
         indole;
    3-(1-benzylpyrrolidin-3-yl)-1-(3,4-dichlorophenyl-sulfonyl)-
         1H-indole;
    3-(1-benzylpyrrolidin-3-yl)-1-(2-bromophenylsulfonyl)-1H-
25
         indole;
    5-[3-(1-benzylpyrrolidin-3-yl)-indole-1-sulfonyl]-4-methyl-
         thiazol-2-ylamine;
    3-(1-benzylpyrrolidin-3-yl)-1-[(5-bromothien-2-yl)sulfonyl]-
30
         1H-indole;
    1-phenylsulfonyl-3-(1-methylpyrrolidin-3-yl)-1H-pyrrolo[2,3-
         b]pyridine;
    1-phenylsulfonyl-3-(1-methylpyrrolidin-3-yl)-1H-indazole;
    1-phenylsulfonyl-3-(1-methyl-2,5-dihydro-1H-pyrrol-3-yl)-1H-
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pyrrolo[2,3-b]pyridine;

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1-phenylsulfonyl-3-(1-methyl-2,5-dihydro-1H-pyrrol-3-yl)-1H-
         indole;
   1-phenylsulfonyl-3-(1-methylpiperidin-4-yl)-1H-indazole;
   1-phenylsulfonyl-3-(1-methyl-1,2,3,6-tetrahydropyridin-4-
         yl)-1H-indazole;
5
   1-phenylsulfonyl-3-(1-methylazepan-4-yl)-1H-pyrrolo[2,3-
         b]pyridine;
    1-phenylsulfony1-3-(1-methylazepan-4-yl)-1H-indole;
    1-phenylsulfonyl-5-fluoro-3-(1-methylazepan-4-yl)-1H-indole;
    1-phenylsulfonyl-3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-
10
         v1)-1H-indole;
    1-phenylsulfonyl-3-(1-methyl-2,5,6,7-tetrahydro-1H-azepin-4-
         yl)-1H-indole;
    1-phenylsulfonyl-3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-
         yl)-1H-pyrrolo[2,3-b]pyridine;
15
    1-phenylsulfonyl-5-fluoro-3-(1-methyl-2,3,6,7-tetrahydro-1H-
         azepin-4-yl)-1H-indole;
    1-phenylsulfonyl-5-fluoro-3-(1-methyl-2,5,6,7-tetrahydro-1H-
         azepin-4-yl)-1H-indole;
    1-(benzo[b]thioen-4-ylsulfonyl)-3-(1-methyl-pyrrolidin-3-
20
         yl)-1H-pyrrolo[2,3-b]pyridine;
    1-(3-fluorophenylsulfonyl)-3-(1-methylpyrrolidin-3-yl)-1H-
         indazole;
    1-(2,5-dichlorophenylsulfonyl)-3-(2,5-dihydro-1H-pyrrol-3-
         v1)-1H-pyrrolo[2,3-b]pyridine;
25
    8-[3-(1-methyl-2,5-dihydro-1H-pyrrol-3-yl)indole-1-
         sulfonyl]-quinoline;
    1-phenylsulfonyl-5-chloro-3-(1-methylpiperidin-4-yl)-1H-
         indazole;
    5-methoxy-3-(1-methyl-1,2,3,6-tetrahydropyridin-4-yl)-1-
30
          (naphth-1-yl-sulfonyl)-1H-indazole;
    3-(1-methylazepan-4-yl)-1-(naphth-1-yl-sulfonyl)-1H-
         pyrrolo[2,3-b]pyridine;
    3-(1-methylazepan-4-yl)-1-(naphth-1-yl-sulfonyl)-1H-indole;
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1-(benzo[b]thien-4-ylsulfonyl)-5-fluoro-3-(1-methylazepan-4-
        yl)-1H-indole;
   8-[3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-yl)-indole-1-
         sulfonyl]-quinoline;
   3-(1-methyl-2,5,6,7-tetrahydro-1H-azepin-4-yl)-1-(naphth-1-
5
         ylsulfonyl)-1H-indole;
    8-[3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-yl)-
         pyrrolo[2,3-b]pyridine-1-sulfonyl]-quinoline;
    8-[5-fluoro-3-(1-methyl-2,3,6,7-tetrahydro-1H-azepin-4-yl)-
         indole-1-sulfonyl]-quinoline;
10
    5-fluoro-3-(1-methyl-2,5,6,7-tetrahydro-1H-azepin-4-yl)-1-
         (naphth-1-ylsulfonyl)-1H-indole;
    1-(benzo[b]thien-4-ylsulfonyl)-3-(1-benzyl-pyrrolidin-3-yl)-
         1H-pyrrolo[2,3-b]pyridine;
    1-(3-fluoro-phenylsulfonyl)-3-(1-phenethyl-pyrrolidin-3-yl)-
15
         1H-indazole;
    1-(2,5-dichlorophenylsulfonyl)-3-(1-ethyl-2,5-dihydro-1H-
         pyrrol-3-yl)-1H-pyrrolo[2,3-b]pyridine;
    3-(1-methyl-2,5-dihydro-1H-pyrrol-3-yl)-1-(naphth-2-
         ylsulfonyl)-1H-indole;
20
    5-chloro-1-(3-fluorophenylsulfonyl)-3-piperidin-4-yl-1H-
         indazole;
    5-methoxy-1-(naphth-1-ylsulfonyl)-3-(1,2,2-trimethyl-
         1,2,3,6-tetrahydro-pyridin-4-yl)-1H-indazole;
    1-(naphth-1-ylsulfonyl)-3-(1-phenethyl-azepan-4-yl)-1H-
25
         pyrrolo[2,3-b]pyridine;
    3-azepan-4-yl-1-(naphth-1-ylsulfonyl)-1H-indole;
    3-azepan-4-yl-1-(3-chloro-5-methyl-benzo[b]thien-2-
         ylsulfonyl)-5-fluoro-1H-indole;
    8-[3-(1-phenethyl-2,3,6,7-tetrahydro-1H-azepin-4-yl)-indole-
30
          1-sulfonyl]-quinoline;
    3-[1-(3,3-dimethylbutyl)-2,5,6,7-tetrahydro-1H-azepin-4-yl]-
          1-(naphth-2-ylsulfonyl)-1H-indole;
    1-(2,3-dichlorophenylsulfonyl)-3-(1-methyl-2,3,6,7-
          tetrahydro-1H-azepin-4-yl)-1H-pyrrolo[2,3-]pyridine;
35
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1-[(3-chloro-5-methoxyphenylsulfonyl)]-3-(2,2-dimethyl-
         2,3,6,7-tetrahydro-1H-azepin-4-yl)-5-fluoro-1H-indole;
   3-azepan-4-yl-5-fluoro-1-(naphth-2-ylsulfonyl)-1H-indole;
    1-Benzenesulfonyl-3-piperidin-3-yl-1H-indole;
   1-(4-isopropyl-benzenesulfonyl)-3-piperidin-3-yl-1H-indole;
5
    1-(5-chloro-thiophene-2-sulfonyl)-3-piperidin-3-yl-1H-
         indole;
    1-(3-chloro-benzenesulfonyl)-3-piperidin-3-yl-1H-indole;
    1-(3,4-difluoro-benzenesulfonyl)-3-piperidin-3-yl-1H-indole;
    1-(4-trifluoromethoxy-benzenesulfonyl)-3-piperidin-3-yl-1H-
10
         indole;
    1-(4-methoxy-benzenesulfonyl)-3-piperidin-3-yl-1H-indole;
    1-(4-trifluoromethy-benzenesulfonyl)-3-piperidin-3-yl-1H-
         indole;
    1-(3-chloro-4-methyl-benzenesulfonyl)-3-piperidin-3-yl-1H-
15
         indole;
    1-(2-chloro-4-trifluoromethyl-benzenesulfonyl)-3-piperidin-
         3-y1-1H-indole;
    1-(2-naphthylenesulfonyl)-3-piperidin-3-yl-1H-indole;
    1-(5-chloro-3-methyl-benzo[b]thiophene-2-sulfonyl)-3-
20
         piperidin-3-yl-1H-indole;
    1-(2,6-dichloro-imidazo[2,1-b]thiazole-5-sulfonyl)-3-
         piperidin-3-yl-1H-indole;
    2-chloro-3-(3-piperidin-3-yl-indole-1-sulfonyl)-imidazo[1,2-
25
         alpyridine;
    2-chloro-3-(3-piperidin-3-yl-indole-1-sulfonyl)-
         benzo[d]imidazo[2,1-b]thiazole;
    1-(4-isopropyl-benzenesulfonyl)-3-piperidin-3-yl-1H-
         pyrrolo[2,3-b]pyridine;
    1-(5-chloro-thiophene-2-sulfonyl)-3-piperidin-3-yl-1H-
30
         pyrrolo[2,3-b]pyridine;
    1-(3-chloro-benzenesulfonyl)-3-piperidin-3-yl-1H-
         pyrrolo[2,3-b]pyridine;
     1-(3,4-difluoro -benzenesulfonyl)-3-piperidin-3-yl-1H-
          pyrrolo[2,3-b]pyridine;
35
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- 1-(4-trifluoromethoxy-benzenesulfonyl)-3-piperidin-3-yl-1Hpyrrolo[2,3-b]pyridine;
- 1-(3-chloro-4-methyl-benzenesulfonyl)-3-piperidin-3-yl-1Hpyrrolo[2,3-b]pyridine;
- 5 1-(2-chloro-4-trifluoromethyl-benzenesulfonyl)-3-piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
 - 1-(2-naphthylenesulfonyl)-3- piperidin-3-yl-1H-pyrrolo[2,3b]pyridine;
 - 1-(5-chloro-3-methyl-benzo[b]thiophene-2-sulfonyl)-3-piperidin-3-yl-1H-pyrrolo[2,3-b]pyridine;
 - 2-chloro-3-(3-piperidin-3-yl-pyrrolo[2,3-b]pyridine-1-sulfonyl)-imidazo[1,2-a]pyridine;
 - 2-chloro-3-(3-piperidin-3-yl-pyrrolo[2,3-b]pyridine-1-sulfonyl)-benzo[d]imidazo[2,1-b]thiazole; and
- 15 the pharmaceutically acceptable salts thereof.
 - 20. A process for the preparation of a compound of formula If

$$R_2$$
 (CR_3R_4)
 R_1
 SO_2R_5
 (If)

10

wherein

W is N or CR₆;

X is N or CR,;

Y is NR_8 or CR_9R_{10} ;

n is 0 or an integer of 1 or 2;

| | Z is NR_{11} or $CR_{12}R_{13}$ with the proviso that when n is 1 and |
|----|---|
| | W is CR_6 then Z must be $CR_{12}R_{13}$ and with the further |
| | provisos that when Y is NR_s then Z must be $CR_{12}R_{13}$ |
| | and at least one of Y and Z must be NR_8 or NR_{11} ; |
| 5 | R_1 , R_2 and R_7 are each independently H, halogen, CN, |
| | OCO_2R_{14} , CO_2R_{15} , $CONR_{29}R_{30}$, $CONR_{29}R_{30}$, $CNR_{16}NR_{17}R_{18}$, SO_mR_{19} , |
| | $NR_{20}R_{21}$, OR_{22} , COR_{23} or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - |
| | C_6 alkynyl, C_3 - C_6 cycloalkyl, cycloheteroalkyl, aryl |
| | or heteroaryl group each optionally substituted; |
| 10 | $\rm R_{_3}$, $\rm R_{_4}$, $\rm R_{_9}$, $\rm R_{_{10}}$, $\rm R_{_{12}}$ and $\rm R_{_{13}}$ are each independently H or an |
| | optionally substituted C ₁ -C ₆ alkyl group; |
| | $\mathrm{R}_{\scriptscriptstyle{5}}$ is an optionally substituted $\mathrm{C}_{\scriptscriptstyle{1}}\text{-}\mathrm{C}_{\scriptscriptstyle{6}}$ alkyl, aryl or |
| | heteroaryl group; |
| | m is 0 or an integer of 1 or 2; |
| 15 | $\mathrm{R_{6}}$ is H, halogen, or an optionally substituted $\mathrm{C_{1}}$ |
| | C_6 alkyl, C_1 - C_6 alkoxy, aryl or heteroaryl group; |
| | $\rm R_{8}$ and $\rm R_{11}$ are each independently H, $\rm CNR_{26}NR_{27}R_{28}$ or a $\rm C_{1}-$ |
| | C_6 alkyl, C_3 - C_6 cycloalkyl, cycloheteralkyl, aryl or |
| | heteroaryl group each optionally substituted; |
| 20 | $R_{14}^{}$, $R_{15}^{}$, $R_{22}^{}$ and $R_{23}^{}$ are each independently H or an |
| | optionally substituted C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2- |
| | C_6 alkynyl, C_3 - C_6 cycloalkyl, cycloheteroalkyl, aryl |
| | or heteroaryl group; |
| | R_{16} , R_{17} , R_{18} , R_{20} , R_{21} , R_{26} , R_{27} , R_{28} , R_{29} and R_{30} are each |
| 25 | independently H or C ₁ -C ₄ alkyl; |
| | R_{19} is an optionally substituted C_1 - C_6 alkyl, aryl or |
| | heteroaryl group; and |
| | represents a single bond or a double bond |
| | which process comprises reacting a compound of formula IVa |

$$R_2$$
 $(CR_3R_4)n$
 N
 N
 H

(IVa)

wherein W, X, Y, Z, n, R_1 , R_2 , R_3 and R_4 are as defined above with a sulfonyl chloride, R_5SO_2Cl , wherein R_5 is defined above in the presence of a base.